FILE 'HOME' ENTERED AT 11:17:57 ON 30 MAR 2007

=> file registry

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.21 0.21

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 11:18:10 ON 30 MAR 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 29 MAR 2007 HIGHEST RN 928707-03-3 DICTIONARY FILE UPDATES: 29 MAR 2007 HIGHEST RN 928707-03-3

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH December 2, 2006

Please note that search-term pricing does apply when conducting SmartSELECT searches.

REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

=>

Uploading C:\Program Files\Stnexp\Queries\10607175_NEW.str

chain nodes :

8 9 10 15 16 17 18 19

ring nodes :

1 2 3 4 5 6

chain bonds :

2-15 5-8 8-9 9-10 15-16 15-19 16-17 16-18

ring bonds :

1-2 1-6 2-3 3-4 4-5 5-6

exact/norm bonds :

5-8 8-9 9-10 15-16 15-19 16-17 16-18

exact bonds :

2-15

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6

isolated ring systems :

containing 1 :

G1:CH2,NH,O,S

G2:OH, SH, COOH, NH2, Cl, Br, I

G3:H,Cy,Hy,Ak

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 8:CLASS 9:CLASS 10:CLASS

15:CLASS

16:CLASS 17:CLASS 18:CLASS 19:CLASS

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR

G1 CH2, NH, O, S

G2 OH,SH,COOH,NH2,Cl,Br,I

G3 H, Cy, Hy, Ak

Structure attributes must be viewed using STN Express query preparation.

=> s l1

SAMPLE SEARCH INITIATED 11:18:27 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 169763 TO ITERATE

1.2% PROCESSED 2000 ITERATIONS INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED) SEARCH TIME: 00.00.01 1 ANSWERS

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**
BATCH **INCOMPLETE**

PROJECTED ITERATIONS:

3371007 TO 3419513

PROJECTED ANSWERS:

1145 TO 2249

L2

1 SEA SSS SAM L1

=> s 11 full

FULL SEARCH INITIATED 11:18:34 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 3378929 TO ITERATE

23.8% PROCESSED 803745 ITERATIONS

1071 ANSWERS

1125 ANSWERS

29.6% PROCESSED 1000000 ITERATIONS INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.28

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**

BATCH **INCOMPLETE**

PROJECTED ITERATIONS:

3378929 TO 3378929

PROJECTED ANSWERS:

3617 TO 3985

L3 1125 SEA SSS FUL L1

=> d scan

L3 1125 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN

IN INDEX NAME NOT YET ASSIGNED

SQL 9

MF C76 H115 N17 O22 S

Absolute stereochemistry.

PAGE 1-B

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):1

L3 1125 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN

IN INDEX NAME NOT YET ASSIGNED

SQL S

MF C63 H102 N14 O22 S

Absolute stereochemistry.

PAGE 1-B

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

=> file medline, caplus, wpids, uspatfull

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

173.00

173.21

FILE 'MEDLINE' ENTERED AT 11:20:14 ON 30 MAR 2007

FILE 'CAPLUS' ENTERED AT 11:20:14 ON 30 MAR 2007 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'WPIDS' ENTERED AT 11:20:14 ON 30 MAR 2007 COPYRIGHT (C) 2007 THE THOMSON CORPORATION

FILE 'USPATFULL' ENTERED AT 11:20:14 ON 30 MAR 2007 CA INDEXING COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

=> s 12

SAMPLE SEARCH INITIATED 11:20:19 FILE 'WPIDS' SAMPLE SCREEN SEARCH COMPLETED - 16401 TO ITERATE

6.1% PROCESSED 1000 ITERATIONS INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED) SEARCH TIME: 00.00.03

2 ANSWERS

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS:

160508 TO 167512

499

PROJECTED ANSWERS:

157 TO

L44 L2

=> d 14 1-4 ibib, abs, hitstr

ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

2004:308396 CAPLUS Full-text

DOCUMENT NUMBER:

140:339072

TITLE:

Preparation of benzamide derivatives as LPA receptor

antagonists

INVENTOR(S):

Terakado, Masahiko; Nakade, Shinji; Seko, Takuya;

Takaoka, Yoshikazu

PATENT ASSIGNEE(S):

Ono Pharmaceutical Co., Ltd., Japan

SOURCE:

PCT Int. Appl., 304 pp.

DOCUMENT TYPE:

CODEN: PIXXD2

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

									APPLICATION NO.								
WO								WO 2003-JP6680									
	W:	ΑE,	AG,	AL,	AM,	AT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,
•		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	ΕE,	ES,	FI,	GB,	GD,	GE,	GH,
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	KR,	ΚZ,	LC,	LK,	LR,	LS,
		LT,	LU,	LV,	ΜÀ,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NI,	NO,	NZ,	OM,	PH,
		PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	TJ,	TM,	TN,	TR,	TT,	TZ,
		UA,	UG,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	ZW						
	RW:						MZ,					ŪĠ,	ZM,	ZW,	AM,	ΑZ,	BY,
		KG,	KZ,	MD,	RU,	ТJ,	TM,	AT,	ΒE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,
		FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	TR,
							CM,										
								AU 2003-241836									
EP	EP 1553075			A1 20050713			EP 2003-733131					20030528					
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,
							RO,										
us									US 2005-530249								
PRIORITY APPLN. INFO.:								JP 2002-291137									
										WO 2	003-	JP66	80		W 2	0030	528
OTHER SOURCE(S):				MAR	PAT	140:	3390										

GΙ

The title compds. I [wherein R = (un) substituted aliphatic hydrocarbyl or cyclyl; G = a bond or a spacer; T = CH2 or a spacer; J = N or CH; B = (un) substituted aliphatic hydrocarbyl or cyclyl; K = a bond or a spacer; Q = a bond or a spacer; ring D = (un) substituted cyclic ring; L = a bond or a spacer; ring E = (un) substituted cyclic ring; n = 0 or 1; M = a bond or a spacer; Z = a acid group] or prodrugs, or salts thereof are prepared as lysophosphatidic acids (LPA) receptor antagonists. For example, the compound II was prepared in a multi-step synthesis. II showed inhibitory activity with IC50 of 0.095 µM against human EDG-2. I are useful for the treatment of urinary diseases, cancer-related diseases, proliferative diseases, inflammatory immune diseases, diseases caused by secretion failures, brain-related diseases, etc. (no data). Formulations containing I as an active ingredient were also described.

IT 679790-63-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of benzamide derivs. as LPA receptor antagonists)

RN 679790-63-7 CAPLUS

CN Benzeneacetic acid, 4-[[[3-(4-chlorophenyl)propyl](3,5-dimethoxybenzoyl)amino]methyl]- (9CI) (CA INDEX NAME)

$$HO_2C - CH_2$$
 $CH_2 - N - C$
 OMe
 OMe

REFERENCE COUNT: 36 THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 2 OF 4 WPIDS COPYRIGHT 2007 THE THOMSON CORP ON STN DATA NOT AVAILABLE FOR THIS ACCESSION NUMBER

L4 ANSWER 3 OF 4 WPIDS COPYRIGHT 2007 THE THOMSON CORP on STN DATA NOT AVAILABLE FOR THIS ACCESSION NUMBER

L4 ANSWER 4 OF 4 USPATFULL on STN

ACCESSION NUMBER:

2006:175380 USPATFULL Full-text

TITLE:

Lpa receptor antagonist

INVENTOR(S):

Terakado, Masahiko, Mishima-gun, JAPAN

Nakade, Shinji, Mishima-gun, JAPAN

Seko, Takuya, Osaka-shi, JAPAN

Takaoka, Yoshikazu, Mishima-gun, JAPAN

PATENT ASSIGNEE(S):

ONO Pharmaceutical Co., LTD. (non-U.S. corporation)

NUMBER KIND DATE

PATENT INFORMATION: US 2006148830 A1 20060706

APPLICATION INFO.: US 2003-530249 A1 20030528 (10)

WO 2003-JP6680 20030528

20050404 PCT 371 date

NUMBER DATE

PRIORITY INFORMATION: JP 2002-291137 20021003

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: SUGHRUE MION, PLLC, 2100 PENNSYLVANIA AVENUE, N.W.,

SUITE 800, WASHINGTON, DC, 20037, US

NUMBER OF CLAIMS: 78
EXEMPLARY CLAIM: 1
LINE COUNT: 7160

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

##STR1## (wherein the symbols are A compound of the general formula (I): as defined in the description), or a non-toxic salt thereof. This compound engages in LPA receptor bonding and antagonism and hence is useful in the prevention and/or treatment of urinary system disease (symptom with prostatic hypertrophy or neurogenic bladder dysfunction disease, symptom to be caused by spinal cord neoplasm, nucleous hernia, spinal canal stenosis or diabetes, occlusion disease of lower urinary tract, inflammatory disease of lower urinary tract, polyuria), carcinoma-associated disease (solid tumor, solid tumor metastasis, angiofibroma, myeloma, multiple myeloma, Kaposi's sarcoma, leucemia and carcinomatous infiltration transition), proliferative disease (disorder with aberrant angiogenesis, artery obstruction and pulmonary fibrosis), inflammation/immune system disease (psoriasis, nephropathy, hepatitis and pneumonitis symptom), disease caused by secretory dysfunction (Sjogren syndrome), brain-related disease (brain infarction, cerebral apoplexy and brain or peripheral neuropathy) or chronic disease (chronic asthma, glomerulonephritis, obesity, prostate hyperplasia, diseases caused by arteriosclerosis process, rheumatism or atopic dermatitis).

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 679790-63-7P

(drug candidate; preparation of benzamide derivs. as LPA receptor antagonists) $\label{eq:candidate}$

RN 679790-63-7 USPATFULL

CN Benzeneacetic acid, 4-[[[3-(4-chlorophenyl)propyl](3,5-dimethoxybenzoyl)amino]methyl]- (9CI) (CA INDEX NAME)

$$HO_2C = CH_2$$
 $CH_2 = N - C$
OMe
OMe

=> d his

(FILE 'HOME' ENTERED AT 11:17:57 ON 30 MAR 2007)

FILE 'REGISTRY' ENTERED AT 11:18:10 ON 30 MAR 2007

L1 STRUCTURE UPLOADED

L2 1 S L1

L3 1125 S L1 FULL

FILE 'MEDLINE, CAPLUS, WPIDS, USPATFULL' ENTERED AT 11:20:14 ON 30 MAR

L4 4 S L2

=>

---Logging off of STN---

=>

Executing the logoff script...

=> LOG Y

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
·	ENTRY	SESSION
FULL ESTIMATED COST	18.02	191.23
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-0.78	-0.78

STN INTERNATIONAL LOGOFF AT 11:21:42 ON 30 MAR 2007